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## AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions and listings of claims in the application:

## LISTING OF CLAIMS:

- 1-18. (Canceled).
- 19. (previously presented): A cell targeting conjugate comprising the following components that are covalently conjugated via a hydrazone, disulphide or amide bond linker that is degradable within the target cells:
- a DNA minor groove binding ligand incorporating an effective Auger electron-emitting, gamma-emitting or positron-emitting atom or photoactive moiety;
  - a target cell specific protein or peptide that is capable of internalisation by target cells; wherein the cell targeting conjugate is represented by Formula (I), wherein:

Formula (I)

X is carbon or nitrogen;

Y1 and Y2 are selected from C(R'), nitrogen, N(R'), oxygen and sulfur, wherein R' is

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hydrogen, optionally substituted alkyl or optionally substituted alkenyl, and wherein  $Y_1$  and  $Y_2$  are not both either C(R') or nitrogen;

 $\stackrel{---}{=}$  is a double bond unless the attached  $Y_1$  or  $Y_2$  is N(R'), oxygen or sulfur in which case it is a single bond;

 $R_1$  to  $R_{12}$  are selected from hydrogen, halogen, hydroxy, amino, optionally substituted alkyl, optionally substituted alkenyl, a moiety including a target cell specific protein or peptide, an Auger electron-emitting moiety, a gamma-emitting moiety, a positron-emitting moiety and a photoactive moiety, and wherein two of  $R_1$  to  $R_5$  may together form optionally substituted cycloalkyl, cycloalkenyl or aryl; wherein one of  $R_1$  to  $R_{12}$  comprises a target cell specific protein or peptide, and wherein one other of  $R_1$  to  $R_{12}$  comprises an Auger electron-emitting moiety, a gamma-emitting moiety, a positron-emitting moiety or a photoactive moiety;

and salts and/or tautomers thereof.

20. (Canceled).

21. (previously presented): The cell targeting conjugate according to claim 19 wherein the target cell specific protein or peptide is selected from anti-A33, C595, 4D5, trastuzumab, egf/R3, humanized h-R3, C225, BrE-3, murine A7, C50, humanized MN-14, anti-A33, MSN-1, bivatuzumab, U36, KIS1, HuM195, anti-CD45, anti-CD19, TXU(anti-CD7)-pokeweed antiviral protein, M195, anti-CD23, apolizumab (Hu1D10), Campath-1H, N901, Ep2, somatostatin analogues, tositumomab, ibritumomab tiuxetan, HB22.7, anti-CD40, OC125, PAM4 and J591.

22-31. (Canceled).

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32. (previously presented) The cell targeting conjugate according to claim 19 wherein the gamma-emitting or positron-emitting moiety is distanced from a DNA minor groove binding region of the conjugate.

- 33. (Canceled).
- 34. (Canceled).
- 35. (Currently amended): A cell targeting conjugate according to claim 32 selected from the following:

$$R_{10} = R_{10} = R$$

and,

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wherein R represents hydrogen, hydroxy, amino, halogen or optionally substituted alkyl, alkenyl or alkynyl, and wherein I represents the gamma-emitting or positron-emitting moiety wherein the gamma-emitting or positron-emitting moiety is iodine (f).